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Ottawa Hull K1A 0C9

(21) (A1) 2,146,707
(22) 1995/04/10
(43) 1995/10/12

(51) Int.Cl. ⁶ C07D 207/34; C07D 307/68; C07D 333/38; C07D 401/00;
C07D 403/02; C07D 405/02; C07D 409/02; A61K 31/33

(19) (CA) APPLICATION FOR CANADIAN PATENT (12)

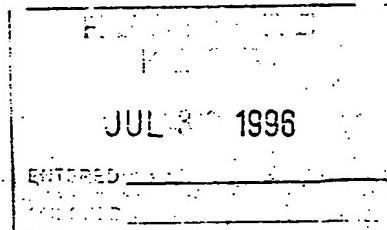
(54) Substituted N-Heteroaroylguanidines, a Process for Their Preparation, Their Use as a Medicament or Diagnostic Agent, and a Medicament Containing Them

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(30) (DE) P 4412334.5 1994/04/11

(57) 17 Claims.



Notice: This application is as filed and may therefore contain an incomplete specification.



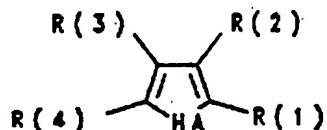
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Abstract

Substituted N-heteroaroylguanidines, a process for their preparation, their use as a medicament or diagnostic agent, and a medicament containing them

The invention relates to heteroaroylguanidines of the Formula I



in which the substituents HA and R(1) to R(5) have the meanings given in claim 1.

These compounds exhibit very good antiarrhythmic properties, as are important for treating diseases which occur, for example, in association with symptoms of oxygen deficiency. As a consequence of their pharmacological properties, the compounds are outstandingly suitable for use as antiarrhythmic pharmaceuticals possessing a cardioprotective component for the prophylaxis and treatment of infarction and for the treatment of angina pectoris, in connection with which they also inhibit or strongly reduce, in a preventive manner, the pathophysiological processes associated with the genesis of ischemically induced damage, in particular associated with the elicitation of ischemically induced cardiac arrhythmias. On account of their protective effects against pathological hypoxic and ischemic situations, the compounds of the formula I according to the invention can, as a consequence of inhibiting the cellular Na^+/H^+ exchange mechanism, be used as pharmaceuticals for treating all acute or chronic damage elicited by ischemia, or diseases induced primarily or secondarily thereby. This is the case with regard to their use as pharmaceuticals for surgical interventions,

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for example in organ transplantations, where the compounds can be used both for protecting the organs in the donor prior to and during removal, for protecting organs which have been removed, for example when they are being treated with or stored in physiological bathing fluids, and when transferring the organs into the recipient. The compounds are likewise valuable protective pharmaceuticals to be used when carrying out angioplastic surgical interventions, for example on the heart or on peripheral vessels. In conformity with their ability to protect against ischemically induced damage, the compounds are also suitable for use as pharmaceuticals for the treatment of ischemias of the nervous system, in particular of the CNS, in connection with which they are suitable, for example, for the treatment of stroke or cerebral edema. Over and above this, the compounds of the formula I according to the invention are also suitable for use in the treatment of forms of shock, such as, for example, allergic, cardiogenic, hypovolemic and bacterial shock.

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Hoechst Aktiengesellschaft

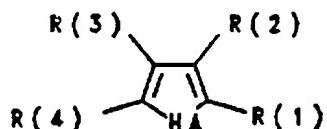
HOE 94/F 094

Dr. v. F.

Description

Substituted N-heteroaroylguanidines, a process for their preparation, their use as a medicament or diagnostic agent, and a medicament containing them

- 5 The invention relates to heteroaroylguanidines of the formula I



in which:

HA is SO_m , O, or NR(5),

m is zero, 1 or 2,

10 R(5) is hydrogen, $(\text{C}_1\text{-}\text{C}_8)$ -alkyl or $-\text{C}_{am}\text{H}_{2am}\text{R}(81)$,

am is zero, 1 or 2

R(81) is $(\text{C}_3\text{-}\text{C}_8)$ -cycloalkyl, or phenyl

which is not substituted or is substituted by 1-3 substituents from the group

15 F, Cl, CF_3 , methyl, methoxy or NR(82)R(83), with R(82) and R(83) being H or CH_3 ;

or

R(81) is $(\text{C}_1\text{-}\text{C}_9)$ -heteroaryl

20 which is linked via C or N and which is unsubstituted or is substituted by 1-3 substituents from the group F, Cl, CF_3 , CH_3 , methoxy, hydroxyl, amino, methyl-amino, or dimethylamino;

25 one of the two substituents R(1) and R(2)

is $-\text{CO-N}=\text{C}(\text{NH}_2)_2$,

and whichever is the other is

hydrogen, F, Cl, Br, I, $(\text{C}_1\text{-}\text{C}_3)$ -alkyl, $-\text{OR}(6)$,

$\text{C}_r\text{F}_{2r+1}$, $-\text{CO-N}=\text{C}(\text{NH}_2)_2$ or $-\text{NR}(6)\text{R}(7)$,

30 R(6) and R(7) are, independently, hydrogen or $(\text{C}_1\text{-}\text{C}_3)$ -alkyl,

r is 1, 2, 3 or 4,

R(3) and R(4) are, independently of each other,
hydrogen, F, Cl, Br, I, -C≡N, X-(CH₂)_p-(C_q-F_{2q+1}),
R(8)-SO_{bm}, R(9)R(10)N-CO, R(11)-CO-

5 or R(12)R(13)N-SO₂-,

where the perfluoroalkyl group is straight-chain
or branched,

X is oxygen, S or NR(14),

R(14) is H or (C₁-C₃)-alkyl,

10 bm is zero, 1 or 2,

p is zero, 1 or 2,

q is zero, 1, 2, 3, 4, 5 or 6,

R(8), R(9), R(11) and R(12) are, independently,
(C₁-C₈)-alkyl, (C₃-C₆)-alkenyl, -C_nH_{2n}-

15 R(15) or CF₃,

n is zero, 1, 2, 3 or 4,

R(15) is (C₃-C₇)-cycloalkyl, or phenyl
which is not substituted or is substituted
by 1-3 substituents from the group
F, Cl, CF₃, methyl, methoxy or NR(16)R(17)
with R(16) and R(17) being H or C₁-C₄-
alkyl.

where R(9), R(11) and R(12) also have the meaning
of H,

25 R(10) and R(13) are, independently,

H or (C₁-C₄)-alkyl,

where R(9) and R(10) and also R(12) and R(13) can
together be 4 or 5 methylene groups, of which one
CH₂ group can be replaced by oxygen, S, NH, N-CH₃
30 or N-benzyl,

or

R(3) and R(4) are, independently of each other,

(C₁-C₈)-alkyl or -C_{al}H_{2al}R(18),

al is zero, 1 or 2,

35 R(18) is (C₃-C₈)-cycloalkyl, or phenyl
which is not substituted or is substituted
by 1-3 substituents from the group
F, Cl, CF₃, methyl, methoxy or
NR(19)R(20), with R(19) and R(20) being H

- 3 -

or CH_3 ;

or

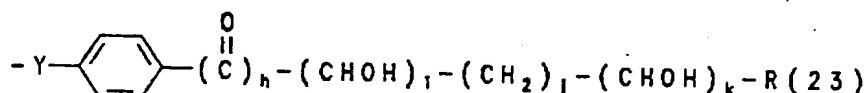
R(3) and R(4) are, independently of each other,
 $(\text{C}_1\text{-}\text{C}_3)$ -heteroaryl,

5

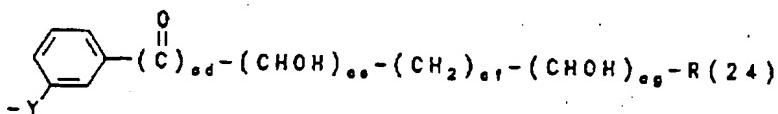
which is linked via C or N and which is unsubstituted or is substituted by 1-3 substituents from the group F, Cl, CF_3 , CH_3 , methoxy, hydroxyl, amino, methylamino or dimethylamino;

or

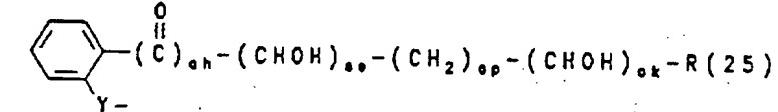
10 R(3) and R(4) are, independently of each other,



or



or



Y is oxygen, -S- or -NR(22)-,
 h , ad and ah are, independently, zero or 1,
 i , j , k , ae , af , ag , ao , ap and ak are,
independently, zero, 1, 2, 3 or 4,

15

where, however, in each case,

h , i and k are not simultaneously zero,
 ad , ae and ag are not simultaneously zero, and
 ah , ao and ak are not simultaneously zero.

R(23), R(24), R(25) and R(22) are, independently,
hydrogen or $(\text{C}_1\text{-}\text{C}_3)$ -alkyl,

20

or

R(3) and R(4) are, independently of each other,
hydrogen, F, Cl, Br, I, CN, $(\text{C}_1\text{-}\text{C}_3)$ -alkyl, $(\text{C}_1\text{-}\text{C}_3)$ -perfluoroalkyl, $(\text{C}_3\text{-}\text{C}_8)$ -alkenyl or $-\text{C}_8\text{H}_{2g}\text{R}(26)$,

25

g is zero, 1, 2, 3 or 4,

- 4 -

R(26) is (C_3-C_8) -cycloalkyl, phenyl, biphenylyl or naphthyl,

5

where the aromatic radicals are not substituted or are substituted by 1-3 substituents from the group F, Cl, CF_3 , methyl, methoxy or $NR(27)R(28)$, with R(27) and R(28) being H, (C_1-C_4) -alkyl or (C_1-C_4) -perfluoroalkyl;

or

10 R(3) and R(4) are, independently of each other,

SR(29), -OR(30), -NR(31)R(32) or -CR(33)R(34)R(35); R(29), R(30), R(31) and R(33) are, independently, $-C_aH_{2a}$ - (C_1-C_9) -heteroaryl

15

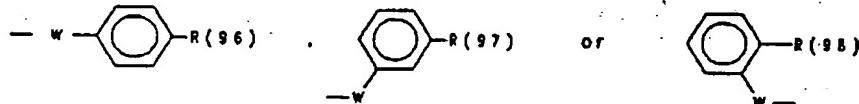
which is unsubstituted or is substituted by 1-3 substituents from the group F, Cl, CF_3 , CH_3 , methoxy, hydroxyl, amino, methylamino or dimethylamino,
a is zero, 1 or 2,

20

R(32), R(34) and R(35) are, independently of each other, defined as R(29), or are hydrogen, (C_1-C_4) -alkyl or (C_1-C_4) -perfluoroalkyl;

or

R(3) and R(4) are, independently of each other,



25

R(96), R(97) and R(98) are, independently, (C_1-C_9) -heteroaryl,

30

which is linked via C or N and which is unsubstituted or is substituted by 1-3 substituents from the group F, Cl, CF_3 , CH_3 , methoxy, hydroxyl, amino, methylamino, dimethylamino or benzyl,

W is oxygen, S or $NR(36)-$,

R(36) is H or (C_1-C_4) -alkyl,

or

R(3) and R(4) are, independently of each other,

- 5 -

R(37) - SO_{cm} or R(38)R(39)N-SO₂-,

cm is 1 or 2,

R(37) is (C₁-C₈)-alkyl, (C₁-C₈)-perfluoroalkyl,
(C₃-C₈)-alkenyl or -C_nH_{2n}-R(40),

5 s is zero, 1, 2, 3 or 4,

R(40) is (C₃-C₈)-cycloalkyl, phenyl, biphenylyl
or naphthyl,

10 where the aromatic radicals are not substituted or are substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(41)R(42), with R(41) and R(42) being H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl;

15 R(38) is H, (C₁-C₈)-alkyl, (C₁-C₈)-perfluoroalkyl,

(C₃-C₈)-alkenyl or -C_wH_{2w}-R(43),

w is zero, 1, 2, 3 or 4,

20 R(43) is (C₃-C₈)-cycloalkyl, phenyl, biphenylyl or naphthyl where the aromatic radicals are not substituted or are substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(44)R(45), with R(44) and R(45) being H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl.

25 R(39) is H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl,

30 where R(38) and R(39) can together be 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl;

or

35 R(3) and R(4) are, independently of each other,

R(46)X(1)-,

35 X(1) is oxygen, S, NR(47), (D=O)A- or NR(48)C=MN(⁺)R(49)-,

M is oxygen or S,

A is oxygen or NR(50),

D is C or SO,

- 6 -

R(46) is (C_1-C_8) -alkyl, (C_3-C_8) -alkenyl,
 $(CH_2)_bC_dF_{2d+1}$ or $-C_xH_{2x}-R(51)$,

b is zero or 1,

d is 1, 2, 3, 4, 5, 6 or 7,

5 x is zero, 1, 2, 3 or 4,

R(51) is (C_3-C_8) -cycloalkyl, phenyl,
biphenylyl, naphthyl,

where the aromatic radicals are not
substituted or are substituted by
1-3 substituents from the group F,
Cl, CF_3 , methyl, methoxy or
NR(52)R(53); with R(52) and
R(53) being H, (C_1-C_4) -alkyl or
 (C_1-C_4) -perfluoroalkyl;

15 R(47), R(48) and R(50) are, independently,
hydrogen, (C_1-C_4) -alkyl or (C_1-C_4) -
perfluoroalkyl,

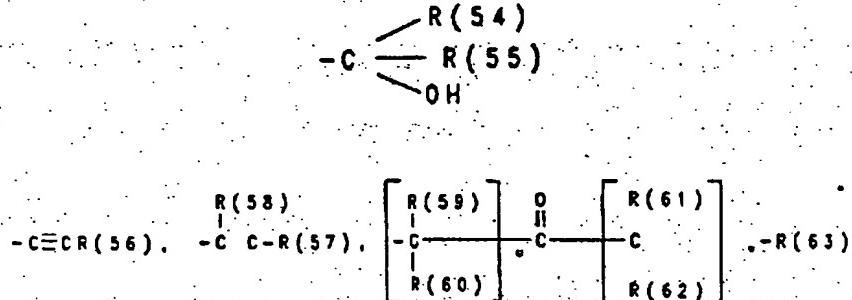
R(49) is defined as R(46), where

20 R(46) and R(47) and, respectively, R(46) and R(48)
can together be 4 or 5 methylene groups, of which
one CH_2 group can be replaced by oxygen, S, NH,
N-CH₃ or N-benzyl,

where A and N^(*) are bonded to the phenyl nucleus
of the benzoylguanidine parent substance;

25 or

R(3) and R(4) are, independently of each other,
-SR(64), -OR(65), -NHR(66), -NR(67)R(68),
-CHR(69)R(70).



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R(64), R(65), R(66), R(67) and R(69) are, identically or differently,

- $(CH_2)_y$ - $(CHOH)_z$ - $(CH_2)_{aa}$ - $(CH_2OH)_t$ -R(71) or
- $(CH_2)_{ab}$ -O- $(CH_2-CH_2O)_{ac}$ -R(72).

5 R(71) and R(72) are hydrogen or methyl,

u is 1, 2, 3 or 4,

v is zero, 1, 2, 3 or 4,

y, z and aa are, identically or differently,

zero, 1, 2, 3 or 4,

10 t is 1, 2, 3 or 4,

R(68), R(70), R(54) and R(55) are, identically or differently,

hydrogen or (C_1-C_6) -alkyl, or

15 R(69) and R(70) and, respectively, R(54) and R(55) are, together with the carbon atom carrying them, a (C_3-C_8) -cycloalkyl;

R(63) is

H, (C_1-C_6) -alkyl, (C_3-C_8) -cycloalkyl or
 $-C_eH_{2e}$ -R(73),

20 e is zero, 1, 2, 3 or 4,

R(56), R(57) and R(73) are, independently,
phenyl,

which is unsubstituted or is substituted by 1-3 substituents from the group F, Cl, CF_3 , methyl,
methoxy or $NR(74)R(75)$ with R(74) and R(75)
being H or (C_1-C_4) -alkyl,

25 or R(56), R(57) and R(73) are, independently,
 (C_1-C_9) -heteroaryl,

which is unsubstituted or is substituted as
phenyl,

30 R(58), R(59), R(60), R(61) and R(62) are hydrogen or
methyl,

or

R(3) and R(4) are, independently of each other,

35 R(76)-NH-SO₂-.

R(76) is R(77)R(78)N-(C=Y')-,

Y' is oxygen, S or N-R(79),

R(77) and R(78) are, identically or
differently,

- 8 -

H, (C_1-C_8)-alkyl, (C_3-C_6)-alkenyl,
or $C_fH_{2f}-R(80)$,

f is zero, 1, 2, 3 or 4,

R(80) is (C_5-C_7)-cycloalkyl, or
phenyl

5

which is unsubstituted or is substituted by 1-3 substituents from
the group F, Cl, CF_3 , methoxy or
(C_1-C_4)-alkyl, or

10 R(77) and R(78) together form 4 or 5 methylene
groups, of which one CH_2 group can be replaced by
oxygen, S, NH, N- CH_3 or N-benzyl, where
R(79) is defined as R(77) or is amidine;

or

15 R(3) and R(4) are, independently of each other,
NR(84)R(85),

R(84) and R(85) are, independently of each other,

H or (C_1-C_4)-alkyl, or, together, can be 4 or 5
methylene groups, of which one CH_2 group can be
replaced by oxygen, S, NH, N- CH_3 or N-benzyl,
or of which one or two CH_2 groups can be
replaced by $CH-C_{dm}H_{2dm+1}$.

20

and the pharmaceutically tolerated salts thereof,

25 where, however, compounds are excepted in which the
radicals R(1) to R(4) and also HA are combined in the
following manner:

R(1)	R(2)	R(3)	R(4)	HA
CON=C(NH ₂)	H	H	Et	O
CON=C(NH ₂)	H	H	Me	O
30 CON=C(NH ₂)	H	H	H	O

Compounds of the formula I are preferred in which:

HA is SO_m , O or NR(5),

m. is zero, 1 or 2,

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- R(5) is hydrogen or methyl,
one of the two substituents R(1) and R(2) is
-CO-N=C(NH₂)₂,
and whichever is the other is hydrogen, F, Cl, CH₃, -OH
5 or -CO-N=C(NH₂)₂,
R(3) is hydrogen, F, Cl, Br, I, -C≡N, C_q-F_{2q+1}, R(8)-SO₂,
R(9)R(10)N-CO, R(11)-CO- or R(12)R(13)N-SO₂-,
where the perfluoroalkyl group is straight-chain
or branched,
10 q is zero, 1, 2, 3, 4, 5 or 6,
R(8), R(9), R(11) and R(12) are, independently,
(C₁-C₈)-alkyl, (C₃-C₄)-alkenyl, -C_nH_{2n}-R(15) or
CF₃,
n is zero, 1, 2, 3 or 4,
15 R(15) is (C₃-C₆)-cycloalkyl, or phenyl
which is not substituted or is substi-
tuted by 1 - 2 substituents from the
group F, Cl, CF₃, methyl, methoxy or
NR(16)R(17), with R(16) and R(17) being H
20 or methyl,
where R(9), R(11) and R(12) also have the meaning
of H;
R(10) and R(13) are, independently, H or methyl,
or
25 R(3) is (C₁-C₈)-alkyl or -C_{al}H_{2al}R(18).
al is zero, 1 or 2,
R(18) is (C₃-C₆)-cycloalkyl, or phenyl
which is not substituted or is substi-
tuted by 1 - 2 substituents from the
group F, Cl, CF₃, methyl, methoxy or
NR(19)R(20), with R(19) and R(20) being H
30 or CH₃,
or
R(3) is quinolyl, isoquinolyl, pyrrolyl, pyridyl or
imadazolyl which are linked via C or N and
which are unsubstituted or are substituted by
1 - 2 substituents from the group F, Cl, CF₃,
CH₃, methoxy, hydroxyl, amino, methylamino or
dimethylamino;

- 10 -

or

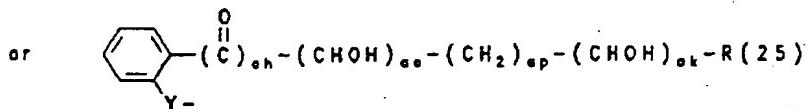
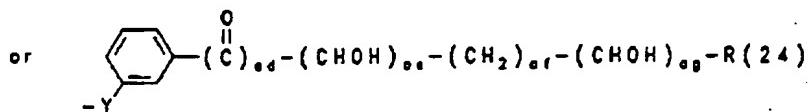
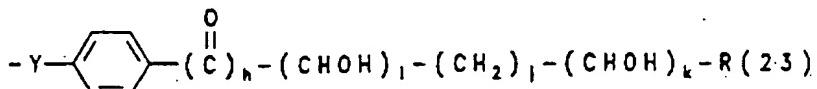
R(3) is $-C\equiv CR(56)$,

R(56) is phenyl,

5

which is unsubstituted or is substituted by
 1 - 2 substituents from the group F, Cl, CF_3 ,
 methyl, methoxy or $NR(16)R(17)$, with R(16) and
 R(17) being H or CH_3 ,

R(4) is



10

 Y is oxygen, $-S-$ or $-NR(22)-$, h , ad and ah are, independently, zero or 1, i , k , ag , ao and ak are, independently, zero, 1, 2 or 3, j , af and ap are, independently, zero or 1,
where, however, in each case,

15

 h , i and k are not simultaneously zero, ad , ae and ag are not simultaneously zero, and ah , ao and ak are not simultaneously zero,R(23), R(24), R(25) and R(22) are, independently,
hydrogen or methyl,

20

or

R(4) is hydrogen, F, Cl, Br, CN, (C_1-C_8) -alkyl, C_q-F_{2q+1} ,
 (C_3-C_8) -alkenyl or $-C_gH_{2g}R(26)$,where the perfluoroalkyl group is straight-chain
or branched,

25

 q is zero, 1, 2, 3 or 4, g is zero, 1 or 2,

- 11 -

R(26) is (C_3-C_8) -cycloalkyl, or phenyl which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF_3 , methyl, methoxy or $NR(27)R(28)$, with R(27) and R(28) being H or CH_3 ,

5
or

R(4) is $SR(29)$, $-OR(30)$, $-NR(31)R(32)$ or $-CR(33)R(34)R(35)$;

10 R(29), R(30), R(31) and R(33) are, independently, $-C_aH_{2a}$ - (C_1-C_9) -heteroaryl, selected from the group consisting of pyrrolyl, imidazolyl, pyrazolyl and pyridyl,

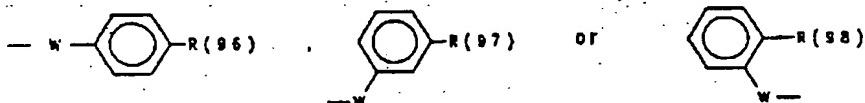
15 which is unsubstituted or is substituted by 1 - 2 substituents from the group F, Cl, CF_3 , CH_3 , methoxy, hydroxyl, amino, methylamino or dimethylamino,
a is zero or 1,

R(32), R(34) and R(35) are, independently of each other,

20 hydrogen or CH_3 ,

or

R(4) is



R(96), R(97) and R(98) are, independently, pyrrolyl, imidazolyl, pyrazolyl or pyridyl,

25 which, in each case, is unsubstituted or is substituted by 1 - 2 radicals from the group comprising

F, Cl, CF_3 , CH_3 , methoxy, dimethylamino or benzyl,

30 W is oxygen, S or $NR(36)-$,

R(36) is H or methyl,

or

R(4) is $R(37)-SO_{cm}$ or $R(38)R(39)N-SO_2-$,

R(37) is (C_1-C_6) -alkyl, CF_3 , (C_3-C_4) -alkenyl or

-C_sH_{2s}-R(40),

s is zero or 1,

R(40) is (C₃-C₆)-cycloalkyl, or phenyl

which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(41)R(42), with R(41) and R(42) being H or CH₃,

R(38) is H, (C₁-C₄)-alkyl, CF₃, (C₃-C₄)-alkenyl or

-C_wH_{2w}-R(43),

w is zero or 1

R(43) is (C₃-C₈)-cycloalkyl, or phenyl

which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(44)R(45), with R(44) and R(45) being H, (C₁-C₄)-alkyl or CH₃,

R(39) is H or CH₃,

where R(38) and R(39) can together be 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl;

or

R(4) is R(46)X(1)-,

X(1) is oxygen, S, NR(47), (C=O)A- or NR(48)C=MN⁽⁺⁾R(49)-,

M is oxygen,

A is oxygen or NR(50),

R(46) is (C₁-C₆)-alkyl, (C₃-C₄)-alkenyl,

(CH₂)_bC_dF_{2d+1} or -C_xH_{2x}-R(51),

b is zero or 1,

d is 1, 2, 3, 4, 5, 6 or 7,

x is zero or 1,

R(51) is (C₃-C₈)-cycloalkyl, or phenyl

which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(52)R(53); with R(52) and R(53) being H or CH₃,

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R(47), R(48) and R(50)

are hydrogen or (C₁-C₄)-alkyl,

R(49) is defined as R(46), where

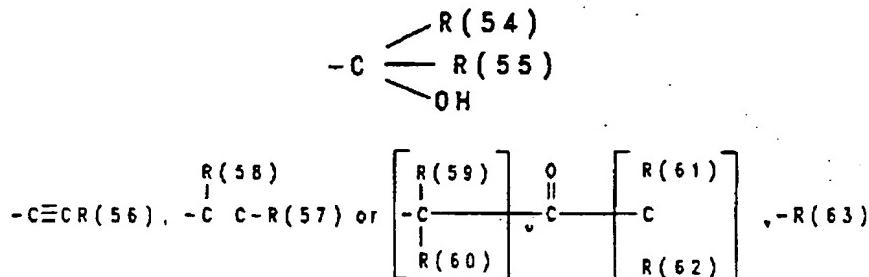
5

R(46) and R(47) and, respectively, R(46) and
R(48) can together be 4 or 5 methylene groups;
of which one CH₂ group can be replaced by
oxygen, S, NH, N-CH₃, or N-benzyl,

10

where A and N^(*) are bonded to the phenyl
nucleus of the benzoylguanidine parent
substance;

or

R(4) is -SR(64), -OR(65), -NHR(66), -NR(67)R(68),
-CHR(69)R(70),

15

R(64), R(65), R(66), R(67) and R(69) are,
identically or differently,

20

-(CH₂)_y-(CHOH)_z-(CH₂)_{aa}-(CH₂OH)_t-R(71) or-(CH₂)_{ab}-O-(CH₂-CH₂O)_{ac}-R(72),

R(71) and R(72) are hydrogen or methyl,

u is 1 or 2,

v is zero, 1 or 2,

y, z and aa are, identically or differently,

zero, 1 or 2,

t is 1, 2 or 3.

25

R(68), R(70), R(54) and R(55) are, identically or
differently,hydrogen or CH₃,

or

R(69) and R(70) and, respectively, R(54) and R(55)
are, together with the carbon atom carrying them, a

(C₃-C₆) -cycloalkyl;

R(63) is

H, (C₁-C₄)-alkyl, (C₃-C₆) -cycloalkyl or
-C_eH_{2e}-R(73),

5 e is zero, 1 or 2,

R(56), R(57) and R(73) are, independently,
phenyl

which is unsubstituted or is substituted
by 1 - 2 substituents from the group F,
10 Cl, CF₃, methyl, methoxy or NR(74)R(75),
with R(74) and R(75) being H or CH₃,

or

R(56), R(57) and R(73) are, independently,

(C₁-C₉)-heteroaryl, selected from the group consist-
15 ing of pyrrolyl, imidazolyl, pyrazolyl and pyridyl,
which is unsubstituted or is substituted
as phenyl;

R(58), R(59), R(60), R(61) and R(62)
are hydrogen or methyl,

20 or

R(4) is R(76)-NH-SO₂-,

R(76) is R(77)R(78)N-(C=Y')-,

Y' is oxygen, S or N-R(79),

25 R(77) and R(78) are, identically or
differently,

H, (C₁-C₄)-alkyl, (C₃-C₄)-alkenyl or
-C_fH_{2f}-R(80),

f is zero or 1,

R(80) is

30 (C₅-C₇)-cycloalkyl, or phenyl

which is unsubstituted or is
substituted by 1 - 2 substi-
tuents from the group F, Cl,
CF₃, methoxy or CH₃, or

35 R(77) and R(78) together form 4 or 5
methylene groups, of which one CH₂ group
can be replaced by oxygen, S, NH, N-CH₃ or
N-benzyl, where

R(79) is defined as R(77).

or

R(4) is NR(84)R(85),

R(84) and R(85) are, independently of each other,

5

H or (C_1-C_4)-alkyl, or together form 4 or 5
methylene groups, of which one CH_2 group can be
replaced by oxygen, S, NH, N- CH_3 or N-benzyl,
or of which one or two CH_2 groups can be
replaced by $CH-CH_3$.

Compounds of the formula I are particularly preferred in
10 which:

R(1) is

-CO-N=C(NH_2)₂

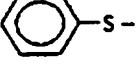
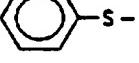
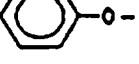
HA is

S, O, NH or N CH_3 ,

15 and the radicals R(2) to R(4) are combined as follows:

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R(2)	R(3)	R(4)
H	n-BuNH-	Cl
H	H ₂ NSO ₂ ⁻	
H	MeSO ₂	
H	 N-	Me
H	 N-	 O-
H	 N-	Me
H	 N-	Cl
H	 N-	MeSO ₂ ⁻

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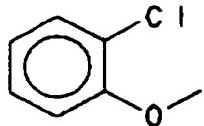
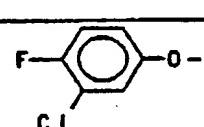
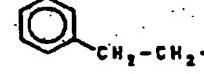
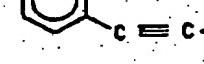
H	MeSO_2	NH_2
H	MeSO_2^-	
H	Cl-	
H	MeSO_2	$(\text{CH}_3)_2\text{CHCH}_2\text{O}-$
H	MeSO_2^-	
H	MeSO_2^-	
H	MeSO_2	

H	Cyclopentyl	2-Chloro-6-methoxybenzene
H	Cyclopentyl	2,6-Dichloro-4-methoxybenzene
H	Cyclopentyl	2-Chloro-6-methylbenzene
H	Cyclopentyl	2-Chloro-4-methoxybenzene
H	Cyclopentyl	2-Methoxy-4-nitrobenzene
H	Cyclopentyl	2-Methoxy-5-nitrobenzene
H	MeSO ₂ ⁻	2-Chloro-4-methylsulfophenyl
H	MeSO ₂ ⁻	2-Chloro-6-methylsulfophenyl
Me	Me	H
H	MeSO ₂ ⁻	i-Pr
H	CF ₃	H

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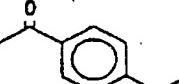
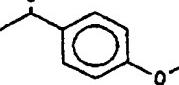
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H		Cl
H	MeSO ₂ -	MeNH-
H	MeSO ₂ -	Et ₂ N-
H	t-Bu	OH
H	MeSO ₂ -	
H	MeSO ₂ -	2-Naphthyl
H	MeSO ₂ -	
H		Me
H		
H	Cl	Et ₂ N-
H	Me ₂ N-	H

H	MeSO ₂ ⁻	
H	Br	NH ₂
H	Cl	H
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
H	CF ₃	CF ₃
H	Me	Me
H	I	CF ₃
H	Me	H
H	H	t-Bu
H	MeSO ₂ ⁻	
H	Me	Cl
H	Br	Me
H	Cl	MeO-
H	MeCO-	
H	Br	Br
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
NH ₂	Br	Me

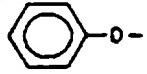
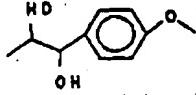
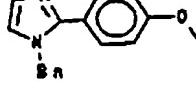
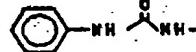
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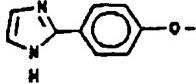
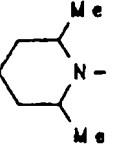
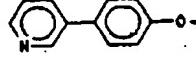
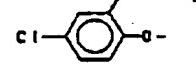
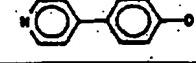
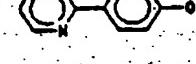
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H	Me ₂ N-	t-Bu
H	MeSO ₂ -	
H		H
H		MeO-
H	Me	Br
H	Cl	F
H	t-Bu	H
NH ₂	Cl	H
H		Me ₂ N
H	Me ₂ N	Cl
H	MeSO ₂ -	7-Isoquinolinoxy
H	MeSO ₂ -	6-quinolinoxy
H	MeSO ₂ -	
H	MeSO ₂ -	
H	MeSO ₂ -	(CH ₃) ₂ CH-CH ₂ -
H	MeSO ₂ -	
H	Me ₂ N-	
H	Me ₂ N-	
H	Me	Me ₂ N-

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H		
H	Me	
H	Cl	i-Pr
H		i-Pr
H	MeSO ₂ -	5-quinolinoxy
H		CF ₃
H	i-Pr	MeSO ₂ -
H	i-Pr	CF ₃
H	H	i-Pr
NH ₂	Br	Br
H	MeSO ₂ -	
H		MeSO ₂ -
H	MeSO ₂ -	
H	Cl	
H	Me ₂ N	i-Pr
H	MeHN-	i-Pr
H	Cl	Cl
H	Me	H ₂ N-
H	Cl	H ₂ N

H	MeSO_2^-	
H	MeSO_2^-	
H	$\text{Me}_2\text{N}-$	i-Pr
CF_3	H	CF_3
H	Br	Me
H	Me	Cl
H	Me_2N	Me
H	CF_3	MeHN-
H	$\text{CH}_3\text{CO-}$	$(\text{CH}_3)_2\text{CH-CH}_2$
H	MeSO_2^-	
H	$\text{CF}_3\text{-O-}$	H
H	Me	Me_2N
H	Cl	$\text{Me}_2\text{N-}$
H	MeSO_2^-	
H	$\text{CH}_3\text{CO-}$	i-Pr
H	Br	BnO-
H	CF_3	Br
H	i-Pr	MeO-
H	MeSO_2^-	
H	MeSO_2^-	

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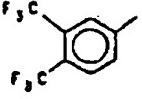
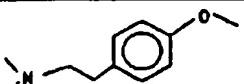
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H	MeO-	t-Bu
H	Br	i-Pr
CF ₃	H	H
H	CF ₃	F
H	Ph	CF ₃
H	CF ₃	1-Imidazolyl
H	MeCO-	t-Butylmethyl
H	Br	F
H	Br	MeO-
H	CF ₃	PhO-
H	CF ₃	Cyclopentyl
H	MesO ₂ -	Cyclobutyl
H	Me	CF ₃
H	MeSO ₂ -	
H	OH	t-Butyl
H	Cl	OMe
H	CF ₃	i-Pr
F	CF ₃	H
F	H	CF ₃
H	t-Butyl	OMe
H	MeCO-	
H	MeCO-	
H	t-Butyl	t-Butyl
H	CF ₃ CF ₂ -	i-Propyl

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H	$\text{CF}_3\text{-SO}_2^-$	
Cl	CF_3	H
Cl	H	CF_3
H	H	Perfluoro-i-propyl
H	H	H
H	MeSO_2^-	
H	H	Perfluoro-n-propyl
H	CF_3	
H	CF_3	
H	CF_3	
H	F	CF_3
H	MeSO_2^-	
H	t-Butyl	i-Propyl
H	t-Butyl	n-Butyl
H	i-Propyl	F
H	i-Butyl	F
H	Cl	1-Imidazolyl
H	H	$\text{CF}_3\text{-CF}_2^-$
H	H	CF_3

H	H	
H	MeSO ₂	
H	CF ₃ SO ₂	i-propyl

(C₁-C₉)-Heteroaryl is understood to mean, in particular, radicals which are derived from phenyl or naphthyl and in which one or more CH groups is/are replaced by N, and/or in which at least two adjacent CH groups are replaced (with the formation of a five-membered aromatic ring) by S, NH or O. In addition, one or both atoms of the condensation site of bicyclic radicals may also be N atoms (as in indolizinyl).

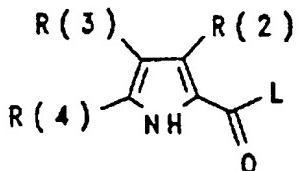
Heteroaryl is, in particular, furanyl, thiienyl, pyrrolyl, imidazolyl, pyrazolyl, triazolyl, tetrazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indolyl, indazolyl, quinolyl, isoquinolyl, phthalazinyl, quinoxalinyl, quinazolinyl and cinnolinyl.

If one of the substituents R(1) to R(5) contains one or more centers of asymmetry, these latter can be either in the S or R configuration. The compounds may be present as optical isomers, as diastereomers, as racemates, or as mixtures thereof.

The designated alkyl radicals may be either straight-chain or branched.

The invention furthermore relates to a process for

preparing compounds I, wherein compounds
of the formula II



in which L is a leaving group which can readily be substituted nucleophilically, are reacted with guanidine.

- 5 The activated acid derivatives of the formula II in which L is an alkoxy, preferably a methoxy, group, a phenoxy group, a phenylthio, methylthio or 2-pyridylthio group, or a nitrogen heterocycle, preferably 1-imidazolyl, are advantageously obtained, in a manner known per se, from the underlying carbonyl chlorides (formula II, L = Cl), which for their part, can be prepared, once again in a manner known per se, from the underlying carboxylic acids (formula II, L = OH), for example using thionyl chloride. In addition to the carbonyl chlorides of the formula II (L = Cl), other activated acid derivatives of the formula II can also be prepared, in a manner known per se, directly from the underlying heteroarylcarboxylic acid derivatives (formula II, L = OH) as can, for example, the methyl esters of the formula II with L = OCH₃, by treatment with gaseous HCl in methanol, the imidazolides of the formula II by treatment with carbonyldiimidazole [L = 1-imidazolyl, Staab, Angew. Chem. Int. Ed. Engl. 1, 351-367 (1962)], the mixed anhydrides II with Cl-COOC₂H₅ or tosyl chloride in the presence of triethylamine in an inert solvent, in addition to which there is also the activation of heteroarylcarboxylic acids with dicyclohexylcarbodiimide (DCC) or with O-[cyano(ethoxy-carbonyl)methylene]amino-1,1,3,3-tetramethyluronium tetrafluoroborate ("TOTU") [Proceedings of the 21st European Peptide Symposium, Peptides 1990, Editors

E. Giralt and D. Andreu, Escom, Leiden, 1991]. A series of suitable methods for preparing activated carboxylic acid derivatives of the formula II is given, with citation of the source literature, on p. 350 in J. March,
5 Advanced Organic Chemistry, Third Edition (John Wiley & Sons, 1985).

An activated carboxylic acid derivative of the formula I is reacted with guanidine, in a manner known per se, in
10 a protic or aprotic polar, but nevertheless inert, organic solvent. In this context, methanol, isopropanol or THF, at a temperature of from 20°C up to the boiling temperature of these solvents, have proved of value when reacting the methyl heteroarylcarboxylates (II, L = OMe) with guanidine. Most of the reactions of compounds II
15 with salt-free guanidine were advantageously carried out in inert solvents such as THF, dimethoxyethane, dioxane or isopropanol. However, water can also be used as the solvent.

When L is Cl, the reaction is advantageously carried out
20 with the addition of an acid-capturing agent, for example in the form of excess guanidine, for binding the hydrohalic acid.

Some of the underlying heteroaryl carboxylic acid derivatives of the formula II are known and described in the literature.
25 The unknown compounds of the formula II may be prepared by methods which are known from the literature, by, for example, converting 5-halo-4-chlorosulfonylbenzoic acids, with ammonia or amines, into 4-amino sulfonyl-5-halo-heteroarylcarboxylic acids, or, with
30 a weak reducing agent, such as sodium bisulfite, and subsequent alkylation, into 4-alkylsulfonyl-5-halo-heteroarylcarboxylic acids, and transforming them, by one of the above-described process variants, into compounds I according to the invention.

35 The introduction of substituted sulfur nucleophiles,

oxygen nucleophiles or nitrogen nucleophiles is achieved using methods, which are known from the literature, for nucleophilic substitution in an aromatic compound. In this substitution, halides and trifluoromethanesulfonates 5 have proved to be of value as leaving groups. The reaction is advantageously carried out in a dipolar aprotic solvent, such as, for example, DMF or TMU, at a temperature of between 0°C and the boiling point of the solvent, preferably between 80°C and the boiling point of the 10 solvent. An alkali metal salt or alkaline earth metal salt having an anion of high basicity and low nucleophilicity, such as, for example, K₂CO₃, is advantageously used as acid-capturing agent.

15 The introduction of the alkyl or aryl substituents is achieved by the methods, which are known from the literature, of palladium-mediated cross-coupling of aryl halides with, for example, organozinc compounds, organostannanes, organoboronic acids or organoboranes.

20 In general, heteroaroylguanidines I are weak bases and can bind acid with the formation of salts. Suitable acid addition salts are the salts of all pharmacologically tolerated acids, for example halides, in particular hydrochlorides, lactates, sulfates, citrates, tartrates, acetates, phosphates, methylsulfonates and p-toluenesulfonates. 25

It was surprising that, while the compounds according to the invention do not exhibit any undesirable and disadvantageous salidiuretic properties, they do exhibit very good antiarrhythmic properties, as are important for 30 treating diseases which occur, for example, in association with symptoms of oxygen deficiency. As a consequence of their pharmacological properties, the compounds are outstandingly suitable for use as antiarrhythmic pharmaceuticals possessing a cardioprotective 35 component for the prophylaxis and treatment of infarction and for the treatment of angina pectoris, in connection

with which they also inhibit or strongly reduce, in a preventive manner, the pathophysiological processes associated with the genesis of ischemically induced damage, in particular associated with the elicitation of
5 ischemically induced cardiac arrhythmias. On account of their protective effects against pathological hypoxic and ischemic situations, the compounds of the formula I according to the invention can, as a consequence of inhibiting the cellular Na^+/H^+ exchange mechanism, be
10 used as pharmaceuticals for treating all acute or chronic damage elicited by ischemia, or diseases induced primarily or secondarily thereby. This is the case with regard to their use as pharmaceuticals for surgical interventions, for example in organ transplantations,
15 where the compounds can be used both for protecting the organs in the donor prior to and during removal, for protecting organs which have been removed, for example when they are being treated with or stored in physiological bathing fluids, and when transferring the
20 organs into the recipient. The compounds are likewise valuable protective pharmaceuticals to be used when carrying out angioplastic surgical interventions, for example on the heart or on peripheral vessels. In conformity with their ability to protect against ischemically induced damage, the compounds are also suitable for
25 use as pharmaceuticals for the treatment of ischemias of the nervous system, in particular of the CNS, in connection with which they are suitable, for example, for the treatment of stroke or cerebral edema. Over and above
30 this, the compounds of the formula I according to the invention are also suitable for use in the treatment of forms of shock, such as, for example, allergic, cardiogenic, hypovolemic and bacterial shock.

In addition to this, the compounds of the formula I according to the invention are notable for their strong inhibitory effect on the proliferation of cells, for example the proliferation of fibroblast cells and the proliferation of the smooth muscle cells of the blood

- vessels. For this reason, the compounds of the formula I are valuable therapeutic agents for use in diseases in which cell proliferation represents a primary or secondary cause and may, therefore, be used as antiatherosclerotic agents, and as agents against diabetic late complications, cancerous diseases, fibrotic diseases such as pulmonary fibrosis, hepatic fibrosis or renal fibrosis, and against organ hypertrophies or hyperplasias, in particular hyperplasia or hypertrophy of the prostate.
- 10 The compounds according to the invention are efficient inhibitors of the cellular sodium/proton antiporter (Na^+/H^+ exchanger), which, in numerous diseases (essential hypertension, atherosclerosis, diabetes, etc.), is also elevated in those cells which are readily accessible to measurement, such as, for example, erythrocytes, thrombocytes or leucocytes. The compounds according to the invention therefore represent outstanding and simple scientific tools, for example in their use as diagnostic agents for defining and differentiating particular forms 15 of hypertension and also of atherosclerosis, diabetes, proliferative diseases, etc. In addition to this, the compounds of the formula I can suitably be used in preventive therapy for preventing the genesis of high blood pressure, for example of essential hypertension.
- 20 25 The compounds according to the invention exhibit a solubility in water which is significantly superior to that of the known compounds. For this reason, their suitability for i.v. administration is considerably greater.
- 30 35 In this context, pharmaceuticals which contain a compound I may be administered orally, parenterally, intravenously or rectally, or by inhalation, the preferred route of administration depending on the given features of the disease. In this context, the compounds I may be used either alone or together with pharmaceutical auxiliary substances, both in veterinary and in human medicine.

Owing to his specialist knowledge, the person skilled in the art is familiar with those auxiliary substances which are suitable for the desired pharmaceutical formulation. Antioxidants, dispersants, emulsifiers, defoamers, taste 5 corrigents, preservatives, solubilizers or dyes, for example, can be used in addition to solvents, gel formers, suppository bases, tablet auxiliaries and other active compound excipients.

For a form for oral use, the active compounds are mixed 10 with the additives, such as carrier substances, stabilizers or inert diluents, which are suitable for the purpose, and brought by the customary methods into the forms, such as tablets, coated tablets, hard gelatin capsules, or aqueous, alcoholic or oily solutions, which 15 are suitable for administration. Gum arabic, magnesium oxide, magnesium carbonate, potassium phosphate, lactose, glucose or starch, in particular corn starch, can, for example, be used as inert excipients. In this context, the preparation can be effected either as a dry granulate 20 or as a wet granulate. Vegetable or animal oils, for example, such as sunflower oil or cod-liver oil, are suitable for use as oily excipients or as solvents.

For subcutaneous or intravenous administration, the active compounds are brought into solution, suspension or 25 emulsion, if desired using the substances, such as solubilizers, emulsifiers or other auxiliary substances, which are customary for the purpose. Examples of suitable solvents are: water, physiological sodium chloride solution or alcohols, for example ethanol, propanol or 30 glycerol, as well as sugar solutions, such as glucose or mannitol solutions, or else a mixture of the different solvents mentioned.

Solutions, suspensions or emulsions of the active compound of the formula I in a pharmaceutically harmless 35 solvent, such as, in particular, ethanol or water, or in a mixture of such solvents, represent examples of

suitable pharmaceutical formulations for administration in the form of aerosols or sprays. As required, the formulation can also contain additional pharmaceutical auxiliary substances such as surfactants, emulsifiers and 5 stabilizers, as well as a propellant gas. Such a preparation customarily contains the active compound in a concentration of from about 0.1 to 10, in particular of from about 0.3 to 3, % by weight.

10 The dosage of the active compound of the formula I to be administered, and the frequency of administration, depend on the strength and duration of the effect of the compounds used; additionally also on the nature and severity of the disease to be treated and on the sex, age, weight and individual responsiveness of the mammalian subject to 15 be treated.

On average, the daily dose of a compound of the formula I is, for a patient of approximately 75 kg in weight, at least 0.001 mg, preferably 0.01 mg to 10 mg, preferably 20 1 mg. In acute manifestations of the disease, for example immediately after suffering a cardiac infarction, even greater and, in particular, more frequent dosages may also be necessary, for example up to 4 individual doses per day. In the case of i.v. use in particular, for example in an infarction patient in intensive care, up to 25 100 mg per day may be necessary.

The novel compounds of the formula I which are listed below, and their physiologically tolerated salts, can be prepared in analogy with the instructions given in the exemplary embodiments:

3.0 List of abbreviations:

MeOH	methanol
DMF	N,N-dimethylformamide
TMU	N,N,N',N'-tetramethylurea
NBS	N-bromosuccinimide

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AIBN	α,α -azobis(isobutyronitrile)
EI	electron impact
DCI	desorption chemical ionisation
RT	room temperature
5 EA	ethyl acetate (EtOAc)
DIP	diisopropyl ether
MTB	methyl tert-butyl ether
mp	melting point
HEP	n-heptane
10 DME	dimethoxyethane
FAB	fast atom bombardment
CH ₂ Cl ₂	dichloromethane
THF	tetrahydrofuran
eq	equivalent
15 ES	electrostatic spray ionization
Me	methyl
Et	ethyl
Bn	benzyl
CNS	central nervous system
20 brine	saturated aqueous solution of NaCl

Experimental Section

Example 1

5-Heptafluoroisopropyl-1-methylpyrrole-2-carboguanidine

a) Methyl 5-heptafluoroisopropyl-1-methylpyrrole-2-carboxylate
 25 1.1 g of methyl 1-methylpyrrole-2-carboxylate, 1.7 ml of perfluoroisopropyl iodide and 1.3 g of FeSO₄ · 7 H₂O are initially introduced in 80 ml of DMSO, and 4.1 ml of H₂O₂ (35%) are slowly added dropwise at RT. The mixture is 30 stirred at RT for 1.5 h and then extracted 3 x with 200 ml of MTB on each occasion, and the organic phase is additionally washed 1 x with 100 ml of water and 2 x with 100 ml of brine. Drying takes place over Na₂SO₄ and the solvent is removed in vacuo. Chromatography using EA/HEP 35 1/4 gives 310 mg of a colorless oil.
 R_f (EA/HEP 1/4) = 0.62 MS (DCI): 308 (M + H)⁺

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b) 5-Heptafluoroisopropyl-1-methylpyrrole-2-carbo-guanidine

310 mg of methyl 5-heptafluoroisopropyl-1-methylpyrrole-2-carboxylate and 295 mg of guanidine are boiled under reflux, for 4 h, in 5 ml of anhydrous isopropanol. The solvent is removed in vacuo and the residue is chromatographed using EA. 123 mg of a colorless oil are obtained.

R_f (EA) = 0.26 MS (ES): 335 ($M + H$)⁺

Conversion into the hydrochloride yields white crystals,
mp 165°C

The title compounds in Examples 2 - 5 are synthesized in analogy with Example 1:

Example 2

5-Heptafluoro-n-propyl-1-methylpyrrole-2-carboguanidine

R_f (EA) = 0.20 MS (ES): 335 ($M + H$)⁺
mp (hydrochloride): 207°C

Example 3

5-Pentafluoroethyl-1-methylpyrrole-2-carboguanidine

R_f (EA) = 0.16 MS (DCI) : 285 ($M + H$)⁺
mp (hydrochloride): 210°C

Example 4

5-Trifluoromethyl-1-methylpyrrole-2-carboguanidine

R_f (EA) = 0.16 MS (DCI): 235 ($M + H$)⁺
mp (hydrochloride): 230°C

Example 5

1-Methylpyrrole-2-carboguanidine

R_f (EA/MeOH 10:1) = 0.13 MS (ES): 167 ($M + H$)⁺
mp (hydrochloride): 255°C

Example 6

5-Isopropyl-4-methylsulfonylthiophene-2-carboguanidine

a) 5-bromothiophene-2-carboxylic acid

10 g of thiophene-2-carboxylic acid are dissolved in

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100 ml of acetic acid and 100 ml of water, and a solution
of 4 ml of bromine in 50 ml of acetic acid and 50 ml of
water is added dropwise, at 0°C, over a period of one
hour. The mixture is subsequently stirred at 0°C for 1 h
5 and the product is then filtered off with suction and
recrystallized from water. 4.8 g of colorless crystals
are obtained, mp 140°C

R_f(MTB 2% HOAc) = 0.54 MS(DCI): 207 (M + H)⁺

b) 5-Bromo-4-chlorosulfonylthiophene-2-carboxylic acid
10 37 g of 5-bromothiophene-2-carboxylic acid are dissolved,
at RT, in 133 ml of chlorosulfonic acid, and this mixture
is stirred at 100°C for 45 min. The mixture is
subsequently poured onto 1 kg of ice and the product is
filtered off with suction. 53 g of a colorless solid are
15 obtained, mp 96°C

R_f(MTB 2% HOAc) = 0.3 MS(DCI): 305 (M + H)⁺

c) 5-Bromo-4-hydroxysulfinylthiophene-2-carboxylic acid
20 27.5 g of sodium sulfite are dissolved in 300 ml of
water, and a total of 35 g of 5-bromo-4-chlorosulfonyl-
thiophene-2-carboxylic acid is added, in portions, at
70°C, with a pH of 9 - 11 being maintained using 10 N
NaOH. The mixture is subsequently stirred at 70°C for 2 h
and then adjusted to pH = 1 with HCl, after which the
product is filtered off with suction. 41 g of colorless
25 crystals are obtained.

mp 195°C (decomposition)

d) 5-Bromo-4-hydroxysulfinylthiophene-2-carboxylic acid,
disodium salt
30 41 g of 5-bromo-4-hydroxysulfinylthiophene-2-carboxylic
acid are suspended in 150 ml of water, and 90 ml of 2 N
NaOH are added (pH = 10). The water is removed in vacuo,
the residue is stirred up in 1 l of acetone, and the
product is filtered off with suction. 46 g are obtained
35 of a colorless, amorphous solid, which is immediately
subjected to further reaction.

e) Methyl 5-bromo-4-methylsulfonylthiophene-2-carboxylate
46 g of the title compound 6 d) are suspended in 150 ml
of DMF, and 32 ml of methyl iodide are added. The mixture
is stirred at 50°C for 5 h and then poured onto 1 l of
5 water; the product is filtered off with suction. 35 g of
a colorless solid are obtained, mp 135°C
 R_f (DIP) = 0.20 MS(DCI): 299 ($M + H$)⁺

f) Methyl 5-isopropyl-4-methylsulfonylthiophene-2-car-
boxylate
10 30 ml of a 2 M solution of isopropylmagnesium chloride in
THF are added to 140 ml of a 0.5 M solution of zinc
chloride in THF. The mixture is stirred at 50°C for 5 h
and the resulting isopropylzinc derivative undergoes
further use as solution A. 6 g of methyl 5-bromo-4-
15 methylsulfonylthiophene-2-carboxylate, 0.6 g of [1,1'-
bis(diphenylphosphino)ferrocene]Pd(II)Cl₂ × CH₂Cl₂, and
180 mg of CuI are stirred, at RT for 10 min, in 100 ml of
anhydrous THF, and solution A is subsequently added
dropwise. The mixture is subsequently stirred at RT for
20 18 h, and the solvent is then removed in vacuo. The
residue is suspended in 200 ml of a saturated aqueous
solution of NaHSO₄, and this suspension is extracted 3 x
with 200 ml of EA on each occasion. Drying takes place
over Na₂SO₄, the solvent is removed in vacuo, and the
25 residue is chromatographed using once in each case, DIP
and EA/HEP 1:3. 1.7 g of a colorless oil are obtained.

R_f (DIP) = 0.29 R_f (EA/HEP 1:3) = 0.32
MS(DCI) : 263 ($M + H$)⁺

g) 5-Isopropyl-4-methylsulfonylthiophene-2-carboguanidine
30 700 mg of methyl 5-isopropyl-4-methylsulfonylthiophene-2-
carboxylate and 790 mg of guanidine are dissolved in 5 ml
of anhydrous isopropanol, and this mixture is boiled
under reflux for 1 h. The solvent is removed in vacuo and
35 80 ml of water are added; the mixture is adjusted to
pH = 2 with aqueous HCl, and the product is filtered off.
The precipitate is dissolved in 50 ml of a saturated
aqueous solution of Na₂CO₃, and this solution is

extracted 3 x with 50 ml of EA on each occasion. The organic phase is dried over Na_2SO_4 , and the solvent is removed in vacuo. 850 mg of an amorphous solid are obtained.

5 R_f (MeOH/EA 1:10) = 0.41 MS(ES) : 290 ($M + H$)⁺
mp (hydrochloride) : 267°C
mp (methanesulfonate) : 128°C

The title compounds of Examples 7, 8 and 10 were synthesized in analogy with Example 6 g):

10 Example 7
5-Methylthiophene-2-carboguanidine
mp (hydrochloride) : 236°C MS(DCI) : 184 ($M + H$)⁺

Example 8
4,5-Dibromo thiophene-2-carboguanidine
15 mp (hydrochloride) : 268°C MS(DCI) : 326 ($M + H$)⁺

Example 9
4-Isopropyl-5-methylsulfonylthiophene-2-carboguanidine
a) 4-Bromo-5-methylthiophene-2-carboxylic acid
25 g of 4,5-dibromo thiophene carboxylic acid, 12.2 g of
20 NaSCH_3 and 60 g of K_2CO_3 are stirred, at 120°C for 5 h, in
1 l of DMF. This mixture is then poured onto 3 l of
water, and the pH of the resulting mixture is adjusted to
1 with HCl; the product is filtered off with suction and
used for further reaction without purification.

25 Yield: 14 g of amorphous powder.

R_f (DIP 2% HOAc) = 0.46

b) 4-Bromo-5-methylsulfonylthiophene-2-carboxylic acid
14 g of the methylthio compound 9 a) are dissolved in
30 500 ml of CH_2Cl_2 , and 41 g of m-chloroperbenzoic acid are
then added in portions. The mixture is stirred at RT for
1.5 h, and the solvent is then removed in vacuo and the
product is esterified without purification.

R_f (DIP 2% HOAc) = 0.10

c) Methyl 4-bromo-5-methylsulfonylthiophene-2-carboxylate
50 ml of SOCl_2 are added to the whole of the crude product from Example 9 b) in 200 ml of MeOH, and this mixture is boiled under reflux for 5 h. Excess SOCl_2 and the solvent are removed in vacuo and the residue is chromatographed using DIP. 11 g of a colorless oil are obtained.

5 R_f (DIP) = 0.28 MS(DCI) : 299 ($M + H$)⁺

d) Methyl 4-isopropyl-5-methylsulfonylthiophene-2-carboxylate
10 30 ml of a 2 M solution of isopropylmagnesium chloride in diethyl ether are added dropwise to a 1 M solution of ZnCl_2 in diethyl ether, and this mixture is boiled under reflux for 6 h. (Solution A)

15 6 g of the bromide 9 c), 588 mg of [1,1-bis(diphenylphosphino)ferrocene]Pd(II)Cl₂ and 183 mg of CuI are stirred, at RT for 10 min, in 100 ml of THF, and solution A is then added to this mixture. The resulting mixture is stirred at RT for 19 h, and 200 ml of EA are then added; 20 the resulting mixture is washed 1 x with 200 ml of water and 1 x with 200 ml of brine. The solvent is removed in vacuo and the residue is chromatographed using EA/HEP 1:2.

2 g of a colorless oil are obtained.

25 R_f (EA/HEP 1:2) = 0.25 MS(DCI) : 263 ($M + H$)⁺

e) 4-Isopropyl-5-methylsulfonylthiophene-2-carboguanidine
1 g of the methyl ester 9 d) is reacted with 1.1 g of guanidine in analogy with Example 6 g). 900 mg of an amorphous powder are obtained.

30 R_f (EA/MeOH 10:1) = 0.41 MS(ES) : 290 ($M + H$)⁺

The compound is converted into the methanesulfonate, mp = 210°C

Example 10

3-Methylthiophene-2-carboguanidine

- 40 -

mp (hydrochloride) : 232°C MS(DCI) : 184 (M + H)⁺**Pharmacological data:****Inhibition of the Na⁺/H⁺ exchanger of rabbit erythrocytes**

- 5 New Zealand White rabbits (Ivanovas) were given a standard diet containing 2% cholesterol for six weeks in order to activate Na⁺/H⁺ exchange and thus to be able to use flame photometry to determine the Na⁺ influx into the erythrocytes via Na⁺/H⁺ exchange. The blood was removed from the aural arteries and rendered incoagulable by the 10 addition of 25 IU of potassium heparin. One part of each sample was used for the duplicate determination of the hematocrit by centrifugation. Aliquots of in each case 100 µl were employed for measuring the initial content of Na⁺ in the erythrocytes.
- 15 In order to determine the amiloride-sensitive sodium influx, 100 µl of each blood sample were in each case incubated, at pH 7.4 and 37°C, in 5 ml of a hyperosmolar salt/sucrose medium (mmol/l: 140 NaCl, 3 KCl, 150 sucrose, 0.1 ouabain, 20 tris(hydroxymethyl)aminomethane). 20 The erythrocytes were then washed three times with ice cold MgCl₂/ouabain solution (mmol/l: 112 MgCl₂, 0.1 ouabain) and hemolyzed in 2.0 ml of distilled water. The intracellular content of sodium was determined by flame photometry.
- 25 The nett influx of Na⁺ was calculated from the difference between the initial sodium values and the sodium content of the erythrocytes following incubation. The amiloride-inhibitable sodium influx was given by the difference in the sodium content of the erythrocytes following incubation with and without 3 × 10⁻⁴ mol/l amiloride. The same procedure was also used in the case of the compounds according to the invention.

Results**Inhibition of the Na⁺/H⁺-exchanger:**

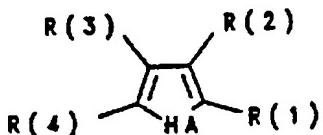
5

10

Example	IC ₅₀ [μmol/l]
1	0.3
2	1.0
3	0.3
4	0.2
5	5.0
6	0.5
7	3
8	0.5

THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE DEFINED AS FOLLOWS:

1. A heteroaroylguanidine of the formula I



in which:

HA is SO_m , O, or NR(5),

5 m is zero, 1 or 2,

R(5) is hydrogen, $(\text{C}_1\text{-C}_8)$ -alkyl or $-\text{C}_{am}\text{H}_{2am}R(81)$,

am is zero, 1 or 2

10 R(81) is $(\text{C}_3\text{-C}_8)$ -cycloalkyl, or phenyl

which is not substituted or is substituted by 1-3 substituents from the group

F, Cl, CF_3 , methyl, methoxy or

NR(82)R(83), with R(82) and R(83) being H

or CH_3 ;

or

15 R(81) is $(\text{C}_1\text{-C}_9)$ -heteroaryl

which is linked via C or N and which is

unsubstituted or is substituted by 1-3

substituents from the group F, Cl, CF_3 ,
CH₃, methoxy, hydroxyl, amino, methyl-

20 amino, or dimethylamino;

one of the two substituents R(1) and R(2)

is $-\text{CO-N}=\text{C}(\text{NH}_2)_2$,

and whichever is the other is

hydrogen, F, Cl, Br, I, $(\text{C}_1\text{-C}_3)$ -alkyl, $-\text{OR}(6)$,

25 $\text{C}_r\text{F}_{2r+1}$, $-\text{CO-N}=\text{C}(\text{NH}_2)_2$ or $-\text{NR}(6)\text{R}(7)$,

R(6) and R(7) are, independently, hydrogen or

$(\text{C}_1\text{-C}_3)$ -alkyl,

r is 1, 2, 3 or 4,

20 R(3) and R(4) are, independently of each other,

hydrogen, F, Cl, Br, I, $-\text{C}\equiv\text{N}$, $\text{X}-\text{(CH}_2\text{)}_p-\text{(C}_q\text{-F}_{2q+1}\text{)}$,

R(8)- SO_{bm} , R(9)R(10)N-CO, R(11)-CO- or R(12)R(13)N-

SO_2^- ,

where the perfluoroalkyl group is straight-chain

or branched,

X is oxygen, S or NR(14),

R(14) is H or (C₁-C₃)-alkyl,

bm is zero, 1 or 2,

5 p is zero, 1 or 2,

q is zero, 1, 2, 3, 4, 5 or 6,

R(8), R(9), R(11) and R(12) are, independently,

(C₁-C₈)-alkyl, (C₃-C₆)-alkenyl, -C_nH_{2n}-

R(15) or CF₃,

10 n is zero, 1, 2, 3 or 4,

R(15) is (C₃-C₇)-cycloalkyl, or phenyl which is not substituted or is substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(16)R(17)

15 with R(16) and R(17) being H or C₁-C₄-alkyl,

where R(9), R(11) and R(12) also have the meaning of H,

R(10) and R(13) are, independently,

20 H or (C₁-C₄)-alkyl,

where R(9) and R(10) and also R(12) and R(13) can together be 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl,

25 or

R(3) and R(4) are, independently of each other,

(C₁-C₈)-alkyl or -C_{al}H_{2al}R(18),

al is zero, 1 or 2,

30 R(18) is (C₃-C₈)-cycloalkyl, or phenyl

which is not substituted or is substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(19)R(20), with R(19) and R(20) being H or CH₃,

35 or

R(3) and R(4) are, independently of each other,

(C₁-C₉)-heteroaryl,

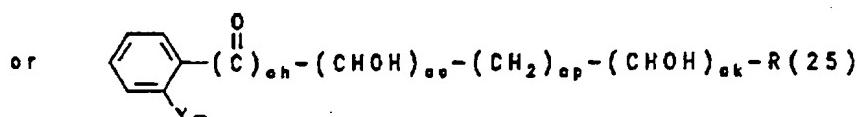
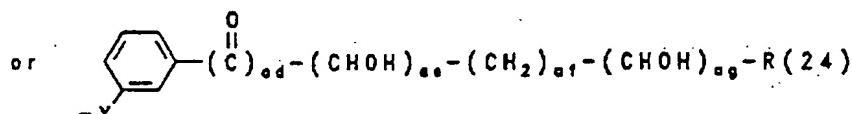
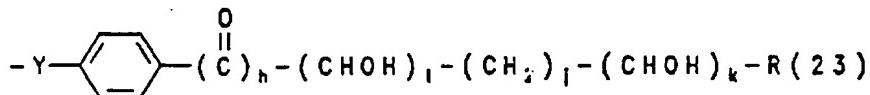
which is linked via C or N and which is unsubstituted or is substituted by 1-3 substituents from

- 44 -

the group F, Cl, CF₃, CH₃, methoxy, hydroxyl, amino, methylamino or dimethylamino;

or

R(3) and R(4) are, independently of each other,



5 Y is oxygen, -S- or -NR(22)-,
 h, ad and ah are, independently, zero or 1,
 i, j, k, ae, af, ag, ao, ap and ak are,
 independently, zero, 1, 2, 3 or 4,
 where, however, in each case,

10 h, i and k are not simultaneously zero,
 ad, ae and ag are not simultaneously zero, and
 ah, ao and ak are not simultaneously zero,
 R(23), R(24), R(25) and R(22) are, independently,
 hydrogen or (C₁-C₃)-alkyl,

15 or

R(3) and R(4) are, independently of each other,
 hydrogen, F, Cl, Br, I, CN, (C₁-C₈)-alkyl, (C₁-C₈)-perfluoroalkyl, (C₃-C₈)-alkenyl or -C₈H_{2g}R(26),

g is zero, 1, 2, 3 or 4,

20 R(26) is (C₃-C₈)-cycloalkyl, phenyl, biphenylyl or naphthyl,
 where the aromatic radicals are not substituted or are substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(27)R(28), with

25

- 45 -

R(27) and R(28) being H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl;

or

R(3) and R(4) are, independently of each other,

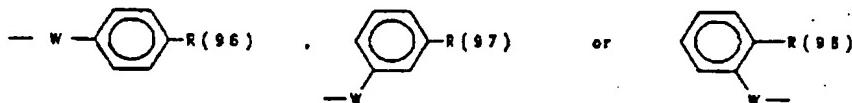
5 SR(29), -OR(30), -NR(31)R(32) or -CR(33)R(34)R(35);
R(29), R(30), R(31) and R(33) are, independently,
-C_aH_{2a}-(C₁-C₉)-heteroaryl

10 which is unsubstituted or is substituted
by 1-3 substituents from the group F, Cl,
CF₃, CH₃, methoxy, hydroxyl, amino,
methylamino or dimethylamino,
a is zero, 1 or 2,

15 R(32), R(34) and R(35) are, independently of each
other, defined as R(29), or are hydrogen, (C₁-
(C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl;

or

R(3) and R(4) are, independently of each other,



R(96), R(97) and R(98) are, independently, (C₁-C₉)-
heteroaryl,

20 which is linked via C or N and which is
unsubstituted or is substituted by 1-3
substituents from the group F, Cl, CF₃,
CH₃, methoxy, hydroxyl, amino, methyl-
amino, dimethylamino or benzyl.

25 W is oxygen, S or NR(36)-,
R(36) is H or (C₁-C₄)-alkyl,

or

R(3) and R(4) are, independently of each other,

R(37)-SO_{cm} or R(38)R(39)N-SO₂-,

30 cm is 1 or 2,
R(37) is (C₁-C₈)-alkyl, (C₁-C₈)-perfluoroalkyl,
(C₃-C₈)-alkenyl or -C_sH_{2s}-R(40),
s is zero, 1, 2, 3 or 4,
R(40) is (C₃-C₈)-cycloalkyl, phenyl, biphenylyl.

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or naphthyl,

where the aromatic radicals are not substituted or are substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(41)R(42), with R(41) and R(42) being H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl;

R(38) is H, (C₁-C₈)-alkyl, (C₁-C₈)-perfluoroalkyl, (C₃-C₈)-alkenyl or -C_wH_{2w}-R(43),

w is zero, 1, 2, 3 or 4,

R(43) is (C₃-C₈)-cycloalkyl, phenyl, biphenylyl or naphthyl where the aromatic radicals are not substituted or are substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(44)R(45), with R(44) and R(45) being H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl,

R(39) is H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl,

where R(38) and R(39) can together be 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl;

25 or

R(3) and R(4) are, independently of each other,

R(46)X(1)-,

X(1) is oxygen, S, NR(47), (D=O)A- or NR(48)C=MN(⁺)R(49)-,

30 M is oxygen or S,

A is oxygen or NR(50),

D is C or SO,

R(46) is (C₁-C₈)-alkyl, (C₃-C₈)-alkenyl, (CH₂)_bC_dF_{2d+1} or -C_xH_{2x}-R(51),

b is zero or 1,

d is 1, 2, 3, 4, 5, 6 or 7,

x is zero, 1, 2, 3 or 4,

R(51) is (C₃-C₈)-cycloalkyl, phenyl, biphenylyl, naphthyl,

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where the aromatic radicals are not substituted or are substituted by 1-3 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(52)R(53); with R(52) and R(53) being H, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl;

5

R(47), R(48) and R(50) are, independently, hydrogen, (C₁-C₄)-alkyl or (C₁-C₄)-perfluoroalkyl;

10

R(49) is defined as R(46), where

15

R(46) and R(47) and, respectively, R(46) and R(48) can together be 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl,

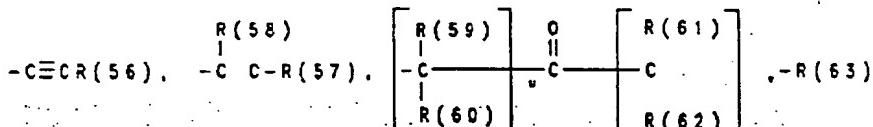
where A and N^(*) are bonded to the phenyl nucleus of the benzoylguanidine parent substance;

or

R(3) and R(4) are, independently of each other,

20

-SR(64), -OR(65), -NHR(66), -NR(67)R(68), -CHR(69)R(70),



R(64), R(65), R(66), R(67) and R(69) are, identically or differently,

25

$-(\text{CH}_2)_y-(\text{CHOH})_z-(\text{CH}_2)_{aa}-(\text{CH}_2\text{OH})_v-\text{R}(71)$ or
 $-(\text{CH}_2)_{ab}-\text{O}-(\text{CH}_2-\text{CH}_2\text{O})_{ac}-\text{R}(72)$,

R(71) and R(72) are hydrogen or methyl,

u is 1, 2, 3 or 4,

v is zero, 1, 2, 3 or 4,

30

y, z and aa are, identically or differently, zero, 1, 2, 3 or 4,

- t is 1, 2, 3 or 4,
R(68), (R70), R(54) and R(55) are, identically or differently,
hydrogen or (C_1-C_6)-alkyl, or
5 R(69) and R(70) and, respectively, R(54) and R(55)
are, together with the carbon atom carrying them, a
(C_3-C_8)-cycloalkyl;
R(63) is
H, (C_1-C_6)-alkyl, (C_3-C_8)-cycloalkyl or
10 - $-C_eH_{2e}-R(73)$,
e is zero, 1, 2, 3 or 4,
R(56), R(57) and R(73) are, independently,
phenyl,
which is unsubstituted or is substituted by 1-3
15 substituents from the group F, Cl, CF_3 , methyl,
methoxy or $NR(74)R(75)$ with R(74) and R(75)
being H or (C_1-C_4)-alkyl,
or R(56), R(57) and R(73) are, independently, (C_1-
 C_9)-heteroaryl,
20 which is unsubstituted or is substituted as
phenyl;
R(58), R(59), R(60), R(61) and R(62) are hydrogen or
methyl,
or
25 R(3) and R(4) are, independently of each other, $R(76)-NH-$
 SO_2^- ,
R(76) is $R(77)R(78)N-(C=Y')-$,
Y' is oxygen, S or N-R(79),
R(77) and R(78) are, identically or
30 differently,
H, (C_1-C_8)-alkyl, (C_3-C_6)-alkenyl,
or $-C_fH_{2f}-R(80)$,
f is zero, 1, 2, 3 or 4,
R(80) is (C_5-C_7)-cycloalkyl, or
phenyl
35 which is unsubstituted or is substituted by 1-3 substituents from
the group F, Cl, CF_3 , methoxy or
(C_1-C_4)-alkyl, or

R(77) and R(78) together form 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl, where R(79) is defined as R(77) or is amidine;

5 or

R(3) and R(4) are, independently of each other,
NR(84)R(85),

R(84) and R(85) are, independently of each other,

10 H or (C₁-C₄)-alkyl, or, together, can be 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl, or of which one or two CH₂ groups can be replaced by CH-C_{dm}H_{2dm+1},

15 and the pharmaceutically tolerated salts thereof,

where, however, compounds are excepted in which the radicals R(1) to R(4) and also HA are combined in the following manner:

	R(1)	R(2)	R(3)	R(4)	HA
20	CON=C(NH ₂)	H	H	Et	O
	CON=C(NH ₂)	H	H	Me	O
	CON=C(NH ₂)	H	H	H	O

2. A heteroaroylguanidine I as claimed in claim 1, wherein:

25 HA is SO_m, O or NR(5),

m is zero, 1 or 2,

R(5) is hydrogen or methyl,

one of the two substituents R(1) and R(2) is
-CO-N=C(NH₂)₂,

30 and whichever is the other is hydrogen, F, Cl, CH₃, -OH or -CO-N=C(NH₂)₂,

R(3) is hydrogen, F, Cl, Br, I, -C≡N, C_q-F_{2q+1},

R(8)-SO₂,

R(9)R(10)N-CO, R(11)-CO- or R(12)R(13)N-SO₂-.

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where the perfluoroalkyl group is straight-chain or branched,

q is zero, 1, 2, 3, 4, 5 or 6,

R(8), R(9), R(11) and R(12) are, independently,

5 (C₁-C₈)-alkyl, (C₃-C₄)-alkenyl, -C_nH_{2n}-R(15) or CF₃,

n is zero, 1, 2, 3 or 4,

R(15) is (C₃-C₆)-cycloalkyl, or phenyl

10 which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(16)R(17), with R(16) and R(17) being H or methyl,

15 where R(9), R(11) and R(12) also have the meaning of H,

R(10) and R(13) are, independently, H or methyl,

or

R(3) is (C₁-C₈)-alkyl or -C_{al}H_{2al}R(18),

20 al is zero, 1 or 2,

R(18) is (C₃-C₆)-cycloalkyl, or phenyl

25 which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(19)R(20), with R(19) and R(20) being H or CH₃;

or

30 R(3) is quinolyl, isoquinolyl, pyrrolyl, pyridyl or imadazolyl which are linked via C or N and which are unsubstituted or are substituted by 1 - 2 substituents from the group F, Cl, CF₃, CH₃, methoxy, hydroxyl, amino, methylamino or dimethylamino;

or

35 R(3) is -C≡CR(56),

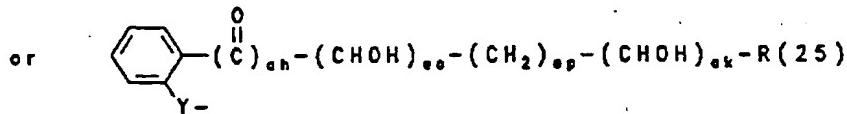
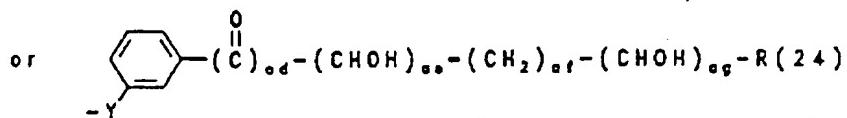
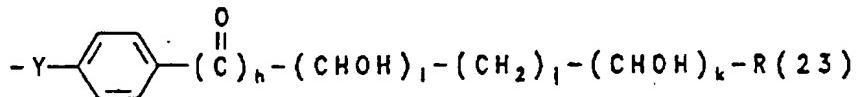
R(56) is phenyl,

which is unsubstituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(16)R(17), with R(16) and

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R(17) being H or CH₃,

R(4) is



Y is oxygen, -S- or -NR(22)-,

h, ad and ah are, independently, zero or 1,

5 i, k, ag, ao and ak are, independently, zero, 1, 2 or 3,

j, af and ap are, independently, zero or 1,

where, however, in each case,

h, i and k are not simultaneously zero,

10 ad, ae and ag are not simultaneously zero, and ah, ao and ak are not simultaneously zero;

R(23), R(24), R(25) and R(22) are, independently, hydrogen or methyl,

or

15 R(4) is hydrogen, F, Cl, Br, CN, (C₁-C₈)-alkyl, C_q-F_{2q+1}, (C₃-C₈)-alkenyl or -C_gH_{2g}R(26),

where the perfluoroalkyl group is straight-chain or branched.

q is zero, 1, 2, 3 or 4,

20 g is zero, 1 or 2,

R(26) is (C₃-C₈)-cycloalkyl, or phenylwhich is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(27)R(28), with R(27) and25 R(28) being H or CH₃.

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or

R(4) is SR(29), -OR(30), -NR(31)R(32) or
-CR(33)R(34)R(35);

5 R(29), R(30), R(31) and R(33) are, independently,
-C_aH_{2a}-(C₁-C₉)-heteroaryl, selected from the group
consisting of pyrrolyl, imidazolyl, pyrazolyl and
pyridyl,

10 which is unsubstituted or is substituted
by 1 - 2 substituents from the group F,
Cl, CF₃, CH₃, methoxy, hydroxyl, amino,
methylamino or dimethylamino,

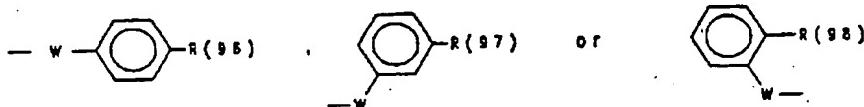
a is zero or 1,

R(32), R(34) and R(35) are, independently of each
other,

15 hydrogen or CH₃,

or

R(4) is



R(96), R(97) and R(98) are, independently, pyrrolyl,
imidazolyl, pyrazolyl or pyridyl;

20 which, in each case, is unsubstituted or is
substituted by 1 - 2 radicals from the group
comprising F, Cl, CF₃, CH₃, methoxy, dimethyl-
amino or benzyl,

W is oxygen, S or NR(36)-,

25 R(36) is H or methyl,

or

R(4) is R(37)-SO_{cm} or R(38)R(39)N-SO₂-,

R(37) is (C₁-C₆)-alkyl, CF₃, (C₃-C₄)-alkenyl or -
C_aH_{2a}-R(40),

30 s is zero or 1,

R(40) is (C₃-C₆)-cycloalkyl, or phenyl
which is not substituted or is substituted
by 1 - 2 substituents from the
group F, Cl, CF₃, methyl, methoxy or

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NR(41)R(42), with R(41) and R(42) being H or CH₃,

R(38) is H, (C₁-C₄)-alkyl, CF₃, (C₃-C₄)-alkenyl or -C_wH_{2w}-R(43),

5 w is zero or 1

R(43) is (C₃-C₈)-cycloalkyl, or phenyl

which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(44)R(45), with R(44) and R(45) being H, (C₁-C₄)-alkyl or CH₃,

10 R(39) is H or CH₃,

15 where R(38) and R(39) can together be 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl;

or

15 R(4) is R(46)X(1)-,

X(1) is oxygen, S, NR(47), (C=O)A- or
20 NR(48)C=MN^(*)R(49)-,

M is oxygen,

A is oxygen or NR(50),

R(46) is (C₁-C₆)-alkyl, (C₃-C₄)-alkenyl,
(CH₂)_bC_dF_{2d+1} or -C_xH_{2x}-R(51),

25 b is zero or 1,

d is 1, 2, 3, 4, 5, 6 or 7,

x is zero or 1,

30 R(51) is (C₃-C₈)-cycloalkyl, or phenyl which is not substituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(52)R(53); with R(52) and R(53) being H or CH₃,

R(47), R(48) and R(50)

35 are hydrogen or (C₁-C₄)-alkyl,

R(49) is defined as R(46), where

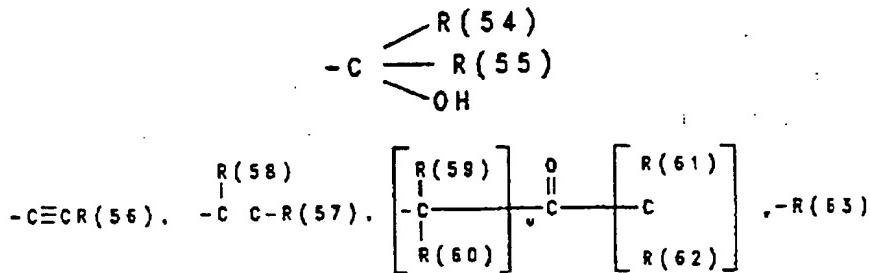
R(46) and R(47) and, respectively, R(46) and R(48) can together be 4 or 5 methylene groups, of which one CH₂ group can be replaced by

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oxygen, S, NH, N-CH₃ or N-benzyl,
where A and N^(*) are bonded to the phenyl
nucleus of the benzoylguanidine parent
substance;

5 or

R(4) is -SR(64), -OR(65), -NHR(66), -NR(67)R(68),
-CHR(69)R(70),



R(64), R(65), R(66), R(67) and R(69) are,
identically or differently,

10 - (CH₂)_y-(CHOH)_z-(CH₂)_{aa}-(CH₂OH)_t-R(71) or
- (CH₂)_{ab}-O-(CH₂-CH₂O)_{ac}-R(72),

R(71) and R(72) are hydrogen or methyl,

u is 1 or 2,

v is zero, 1 or 2,

15 y, z and aa are, identically or differently,
zero, 1 or 2,

t is 1, 2 or 3,

R(68), R(70), R(54) and R(55) are, identically or
differently,

20 hydrogen or CH₃,

or

R(69) and R(70) and, respectively, R(54) and R(55)
are, together with the carbon atom carrying them, a
(C₃-C₆)-cycloalkyl;

25 R(63) is

H, (C₁-C₄)-alkyl, (C₃-C₆)-cycloalkyl or
-C₆H₂e-R(73),

e is zero, 1 or 2,

R(56), R(57) and R(73) are, independently,

30 phenyl

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which is unsubstituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methyl, methoxy or NR(74)R(75), with R(74) and R(75) being H or CH₃.

5 or

R(56), R(57) and R(73) are, independently, (C₁-C₉)-heteroaryl, selected from the group consisting of pyrrolyl, imidazolyl, pyrazolyl and pyridyl,

10 which is unsubstituted or is substituted as phenyl;

R(58), R(59), R(60), R(61) and R(62) are hydrogen or methyl,

or

15 R(4) is R(76)-NH-SO₂-,

R(76) is R(77)R(78)N-(C=Y')-,

Y' is oxygen, S or N-R(79),

R(77) and R(78) are, identically or differently,

20 H, (C₁-C₄)-alkyl, (C₃-C₄)-alkenyl or -C_fH_{2f}-R(80),

f is zero or 1,

R(80) is

(C₅-C₇)-cycloalkyl, or phenyl

25 which is unsubstituted or is substituted by 1 - 2 substituents from the group F, Cl, CF₃, methoxy or CH₃, or

R(77) and R(78) together form 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl, where

R(79) is defined as R(77),

or

35 R(4) is NR(84)R(85),

R(84) and R(85) are, independently of each other,

H or (C₁-C₄)-alkyl, or together form 4 or 5 methylene groups, of which one CH₂ group can be replaced by oxygen, S, NH, N-CH₃ or N-benzyl,

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or of which one or two CH₂ groups can be replaced by CH-CH₃.

3. A heteroaroylguanidine I as claimed in claim 1, wherein:

5 R(1) is

-CO-N=C(NH₂)₂

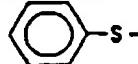
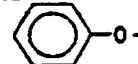
HA is

S, O, NH or NCH₃,

and the radicals R(2) to R(4) are combined as follows:

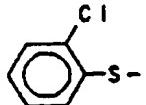
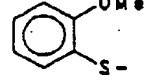
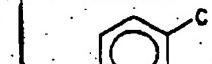
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R(2)	R(3)	R(4)
H	n-BuNH-	Cl
H	H ₂ NSO ₂ ⁻	
H	MeSO ₂	
H		Me
H		
H		Me
H		Cl
H		MeSO ₂ ⁻
H	MeSO ₂	NH ₂
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	

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H	MeSO_2^-	
H	MeSO_2^-	
H	MeSO_2^-	
H	MeSO_2^-	
H	Cl^-	
H	MeSO_2^-	
H	MeSO_2^-	
H	MeSO_2^-	
H	MeSO_2^-	
H	$\text{N}-\text{C}_5\text{H}_9$	

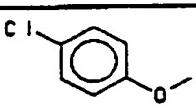
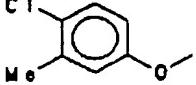
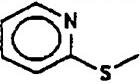
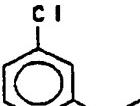
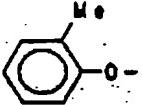
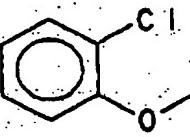
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H	<chem>C1CCN1</chem>	<chem>Oc1ccc(Cl)c(Cl)c1</chem>
H	<chem>C1CCN1</chem>	<chem>Oc1ccc(C)c(Cl)c1</chem>
H	<chem>C1CCN1</chem>	<chem>Oc1ccc(Cl)c(Cl)c1</chem>
H	<chem>C1CCN1</chem>	<chem>Oc1ccc(OC)c(Cl)c1</chem>
H	<chem>C1CCN1</chem>	<chem>Oc1cccnc(Cl)c1</chem>
H	<chem>MSO2^-</chem>	<chem>Oc1ccc(S)c(Cl)c1</chem>
H	<chem>MSO2^-</chem>	<chem>Oc1ccc(S)c(Cl)c1</chem>
Me	Me	H
H	<chem>MSO2^-</chem>	iPr
H	<chem>CF3</chem>	H
H	<chem>C1CCN1</chem>	Cl
H	<chem>MSO2^-</chem>	MeNH-
H	<chem>MSO2^-</chem>	Et ₂ N-

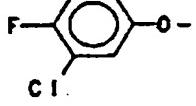
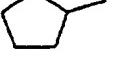
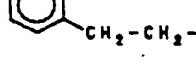
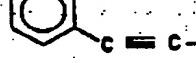
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H	t-Bu	OH
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	2-Naphthyl
H	MeSO ₂ ⁻	
H		Me
H		
H	Cl	Et ₂ N ⁻
H	Me ₂ N-	H
H	MeSO ₂ ⁻	

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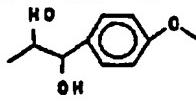
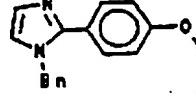
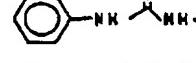
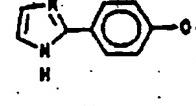
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H	Br	NH ₂
H	Cl	H
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
H	CF ₃	CF ₃
H	Me	Me
H	I	CF ₃
H	Me	H
H	H	t-Bu
H	MeSO ₂ ⁻	
H	Me	Cl
H	Br	Me
H	Cl	MeO-
H	MeCO-	
H	Br	Br
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
NH ₂	Br	Me
H	Me ₂ N-	t-Bu
H	MeSO ₂ ⁻	
H		H

H		MeO-
H	Me	Br
H	Cl	F
H	t-Bu	H
NH ₂	Cl	H
H		Me ₂ N
H	Me ₂ N	Cl
H	MeSO ₂ ⁻	7-Isoquinolinoxy
H	MeSO ₂ ⁻	6-quinolinoxy
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	
H	MeSO ₂ ⁻	(CH ₃) ₂ CH-CH ₂ -
H	MeSO ₂ ⁻	
H	Me ₂ N [.]	
H	Me ₂ N [.]	Cl-
H	Me	Me ₂ N [.]
H		
H	Me	
H	Cl	i-Pr

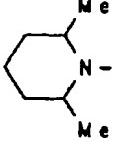
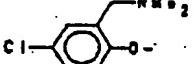
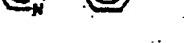
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H		i-Pr
H	MeSO ₂ ⁻	5-quinolinoxy
H		CF ₃
H	i-Pr	MeSO ₂ ⁻
H	i-Pr	CF ₃
H	H	i-Pr
NH ₂	Br	Br
H	MeSO ₂ ⁻	
H		MeSO ₂ ⁻
H	MeSO ₂ ⁻	
H	Cl	
H	Me ₂ N	i-Pr
H	MeHN-	i-Pr
H	Cl	Cl
H	Cl	H ₂ N-
H	Cl	H ₂ N
H	MeSO ₂ ⁻	

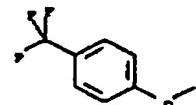
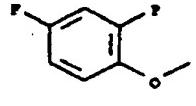
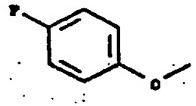
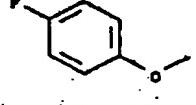
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H	MeSO_2^-	
H	Me_2N^-	i-Pr
CF_3	H	CF_3
H	Br	Me
H	Me	Cl
H	Me_2N	Me
H	CF_3	MeHN^-
H	CH_3CO^-	$(\text{CH}_3)_2\text{CH}-\text{CH}_2$
H	MeSO_2^-	
H	$\text{CF}_3\text{-O}^-$	H
H	Me	Me_2N
H	Cl	Me_2N^-
H	MeSO_2^-	
H	CH_3CO^-	i-Pr
H	Br	BnO^-
H	CF_3	Br
H	i-Pr	MeO^-
H	MeSO_2^-	
H	MeSO_2^-	
H	MeO^-	t-Bu
H	Br	i-Pr
CF_3	H	H
H	CF_3	F

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H	Ph	CF_3
H	CF_3	1-Imidazolyl
H	MeCO-	t-Butylimethyl
H	Br	F
H	Br	MeO-
H	CF_3	PhO-
H	CF_3	Cyclopentyl
H	MeSO_2^-	Cyclobutyl
H	Me	CF_3
H	MeSO_2^-	
H	OH	t-Butyl
H	Cl	OMe
H	CF_3	i-Pr
F	CF_3	H
F	H	CF_3
H	t-Butyl	OMe
H	MeCO-	
H	MeCO-	
H	t-Butyl	i-Butyl
H	CF_3CF_2^-	i-Propyl
H	$\text{CF}_3\text{-SO}_2^-$	
Cl	CF_3	H

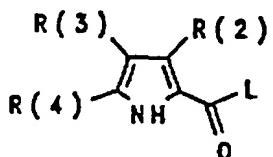
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Cl	H	CF ₃
H	H	Perfluoro-i-propyl
H	H	H
H	MeSO ₂	
H	H	Perfluoro-n-propyl
H	CF ₃	
H	CF ₃	
H	CF ₃	
H	F	CF ₃
H	MeSO ₂ ⁻	
H	t-Butyl	i-Propyl
H	t-Butyl	n-Butyl
H	i-Propyl	F
H	i-Butyl	F
H	Cl	1-Imidazolyl
H	H	CF ₃ -CF ₂ ⁻
H	H	CF ₃
H	H	

H	MeSO ₂	
H	CF ₃ SO ₂	i-Propyl

4. A process for preparing a compound I as claimed in claim 1, wherein a compound of the formula II



in which L is a leaving group which can readily be substituted nucleophilically, is reacted with guanidine.

5. The use of a compound I as claimed in claim 1 for preparing a medicament for the treatment of arrhythmias.

6. A method for treating arrhythmias, wherein an effective quantity of a compound I as claimed in claim 1 is treated with the customary additives and administered in a suitable form for administration.

7. The use of a compound I as claimed in claim 1 for preparing a medicament for the treatment or prophylaxis of cardiac infarction.

15 8. The use of a compound I as claimed in claim 1 for preparing a medicament for the treatment or prophylaxis of angina pectoris.

9. The use of a compound I as claimed in claim 1 for preparing a medicament for the treatment or prophylaxis of ischemic conditions of the heart.

10. The use of a compound I as claimed in claim 1 for preparing a medicament for the treatment or prophylaxis of ischemic conditions of the peripheral and central nervous system and of stroke.

25 11. The use of a compound I as claimed in claim 1 for